AMENDMENTS TO THE CLAIMS

- 1. **(Currently Amended)** A method for the treatment of a tumor which comprises administering to a patient in need thereof an effective amount of active dendritic cells (DC) that are tumor-specific and secrete IL12, said tumor-specific IL12 secreting DC being prepared by a process comprising:
- (a) collecting DC or DC precursor cells from a suitable source to obtain a DC culture;
- (b) loading the DC of said DC culture with a tumor specific antigen; and
- (c) exposing said DC culture to a concentration of LPS and a concentration of IFN-γ effective to trigger the DC of said DC culture to secrete IL12 to thereby obtain said tumor specific and IL12 secreting DC wherein said exposure to LPS and IFN- γ occurs over a period of 1–102-6 hours.
- 2. **(Previously Presented)** The method according to claim 1, wherein said treatment is performed after bone marrow transplantation.
- 3. **(Previously Presented)** The method according to claim 1, wherein said tumor is an advanced malignancy.
- 4. **(Previously Presented)** The method according to claim 1, wherein said DC are collected from the patient having said tumor or from a bone marrow donor.
- 5. **(Previously Presented)** The method according to claim 1, wherein the DCs have been loaded with an antigen from a tumor cell from said patient having said tumor.
- 6. **(Previously Presented)** The method according to claim 5, wherein the DC are additionally charged with a tracer antigen.
- 7. **(Previously Presented)** The method according to claim 6, wherein said tracer antigen is keyhole limpet hemocyanine (KLH).

Docket No.: 4518-0110PUS1

8. **(Previously Presented)** The method according to claim 7, wherein the DCs are additionally charged with an adjuvant, especially with tetanus toxoid.

9. **(Previously Presented)** The method according to claim 1, wherein the DC have been generated in vitro from peripheral blood mononuclear cells (PBMCs).

10.-11. (Cancelled)

- 12. **(Withdrawn)** A method for triggering IL-12 release from dendritic cells (DCs) which comprises administering to a patient an effective amount of a combination of LPS, IFN-γ and a tumor antigen.
- 13. **(Withdrawn)** The method according to claim 12, wherein the DCs have been loaded with an antigen from a tumor cell from a patient having said tumor.
- 14. (Cancelled)
- 15. (Cancelled)
- 16. (Cancelled)
- 17. (Cancelled)
- 18. (Cancelled)
- 19. **(Currently Amended)** A method for the treatment of a tumor which comprises administering to a patient in need thereof an effective amount of active dendritic cells (DC) that are tumor-specific and secrete IL12, said tumor-specific, IL12 secreting DC being prepared by a process consisting essentially of:

- (a) collecting DC or DC precursor cells from a suitable source to obtain a DC culture;
- (b) loading the DC of said DC culture with a tumor specific antigen; and
- (c) exposing said DC culture to a concentration of LPS and a concentration of IFN-γ effective to trigger the DC of said DC culture to secrete IL12 to thereby obtain said tumor specific and IL12 secreting DC wherein said exposure to LPS and IFN- γ occurs over a period of 1–102-6 hours.

20. (Cancelled)

- 21. (Currently Amended) A method for the treatment of a tumor consisting essentially of administering to a patient in need thereof an effective amount of active dendritic cells (DC),[[,]] and wherein said active DC are prepared by a process consisting essentially of:
- (a) collecting DC or DC precursor cells from a suitable source to obtain a DC culture;
- (b) loading the DC of said DC culture with a tumor specific antigen; and exposing said DC culture to a concentration of LPS and a concentration of IFN- γ effective to trigger the DC of said DC culture to secrete IL12 and thereby obtain said active DC wherein said exposure to LPS and IFN- γ occurs over a period of $\frac{1-10}{2}$ -6 hours.
- 22. (**Previously Presented**) The method of claim 1 wherein said active DCs are administered or frozen after exposure to LPS and IFN-γ.
- 23. (**Previously Presented**) The method of claim 1 wherein said active DCs are exposed to LPS and IFN- γ for a period of 2 hours.
- 24. (**Previously Presented**) The method of claim 1 wherein said active DCs are exposed to LPS and IFN- γ for a period of 6 hours.
- 25. (Cancelled)
- 26. (Cancelled)